## CLAINS:

1. A compound of the formula

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wherein A is an L, D or DL amino acid selected from the group consisting of Alanine (Ala), valine (Val), phenylalanine (Phe), para-chloro-phenylalanine (p.Cl Phe), tryptophan (Trp), proline (Pro), serine (Ser), Threonine (Thr), tyrosine (Tyr), glutamic acid (Glu), beta alanine (Beta Ala), -aminobutyric acid (Abu), N-methylalanine (N-Me Ala), 5-fluorotryptopham (5-F Trp), 5-bromotryptopham (5-Br Trp), 5-chlorotryptopham (5-Cl Trp), their acetylated derviatives or a pharmaceutically acceptable acid addition salt thereof;

B is an L, D or DL amino adid selected from the group consisting of threonine amid (Thr NH<sub>2</sub>), valine amide (Val NH<sub>2</sub>), proline amide (Pro NH<sub>2</sub>), hydroxyproline amide (HO Pro NH<sub>2</sub>), serine amide (Ser NH<sub>2</sub>), tyrosine amide (Tyr NH<sub>2</sub>, tryptophan amide (Trp NH<sub>2</sub>), 5-fluorotryptophan amide (5-F Trp NH<sub>2</sub>), formyl tryptophan amide (For Trp NH<sub>2</sub>), alanine amide (Ala NH<sub>2</sub>), glycine amide (Gly NH<sub>2</sub>) and methylalanine amide (Me Ala NH<sub>2</sub>);

X is L-phenylalanine (L-Phe) or L-tyrosine (L-Tyr);

Y is L-threonine (L-Thr) or L-valine (L-Val);

Z is L, D or DL 5-halo-tryptophan, in which the halogen (Halo-) is fluorine, chlorine, bromine or iodine, or D-tryptophan (D-Trp); and

C" and C' are L or D-cysteine (Cys), <- aminobutyric acid (Abu), aspartic acid (Asp) or lysine (Lys);

the connecting line between C" and C' signifies a bridge selected from the group consisting of carbon/carbon, carbon/sulfur, sulfur/sulfur and amide bridges; and the pharmaceutically acceptable acid addition salts thereof.

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A compound according to Claim 1 wherein
                 C" is Cys;
                 X is Phe;
                 Y is Thr; and
                 3. A compound according to Claim \check{\mathcal{X}} wherein
            ि पा C" is Cys;
                X is Tyr;
                Z is D-Trp;
               Y is Val; and
             , 🥴 C' is Cys.
                 4. A compound according to Claim 1 wherein
               C" is Cys;
X is Phe;
                Y is Thr; and
                C' is Cys.
B
                    A compound according to Claim {\mathscr X} wherein
            i di C" is Cys;
               X is Tyr;
                Y is Val; and
              . ⊲C' is Cys.
                6. A compound acoording to Claim 1 wherein
               C" is Lys;
              X is Tyr;
Z is D-Trp;
                Y is Val; and
                 C' is Asp.
                \mathcal{A}. A compound according to Claim \mathcal{X} which is
           D-Phe-Cys-Tyr-D-Trp-Lys-Val-Cys-Thr-NH2.
                 8. A compound according to Claim 1 which is
           Ac-p-Cl-D-Phe-Cys-Phe-D-Trp-Lys-Thr-Cys-Thr-NII2.
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A compound according to Claim which is P.C. - Aus: 4/18/2 OSCI-D-Phe-Cys-Che-D-Trp-Lys-Ch-Cys-Ch-NH2 TOYAOX 10. A compound according to Claim which is TOULT ACOSCI-D-Phe-Cys-Tyr-D-Trp-Lys-Val-Cys-Thr-11112. 1. A compound according to Claim & which is Ac-D-Phe-Lys-Tyr-D-Trp-Lys-Val-Asp-Thr-NH2 (amide bridge). An octapeptide (reduced form) of the formula A - C'' - X - Z - Lys - Y - C' - Ewherein A, C", X, Z, Y, C' and B are as defined in Claim -10' A compound according to Claim which is: D-Phe-Cys-Tyr-D-Trp-Lys-Val-Cys-Thr-NH2. 14. A compound according to Claim 12 which is: D-Phe-Cys-Tyr-D-Trp-Lys-Val-Cys-Ser-NH2. 12
15. A compound according to Claim 12 which is: D-Phe-Cys-Tyr-D-Trp-Lys-Val-Cys-Trp-NH2. R.C. AVS; 4-18. A process for \preparing an octapeptide of formula (I) A - C'' - X - Z - Lys - Y - C' - Bin which A, C", X, Z, Y, C' and B are as defined in Claim 1 which comprises oxidizing a corresponding peptide of formula (II) A - C'' - X - Z - Lys - Y +in which A, C", X, Z, Y, C' and B are as defined above. 17. The process of Claim 16 wherein the peptide of formula (I') is oxidized with potassium ferricyanide, iodine, oxygen or air.

18. A pharmaceutical composition which comprises an octapeptide of Claim 1, its reduced form or a pharmaceutically acceptable acid addition salt thereof in a pharmaceutically acceptable liquid or solid carrier thereof.

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19. A pharmaceutical composition of Claim 18 which comprises an octapeptide of Claim 1, its reduced form or a pharmaceutically acceptable acid addition salt thereof, encapsulated in poly (d,1-lactide-co-glycolide) microcapsules.

20: A method of treating excess release of growth hormone, gastrointestinal disorders, cancer and diabetes in a mammal in need of such therapy which comprises administering to said mammal an effective dose of octapeptide of claim its reduced form, or a pharmaceutically acceptable acid addition salt thereof.

(R.C.-AVP) B) 2+ A COMPOUND ACCORDING TO

(R.C.-AVS)

(R.C.-AVS)

D. Phe-Cys-Tyr-D-Typ-Lys-Val-Cys-Tyr-NH2

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